

Serial Number 07/044,505
Art Unit 1807

The Group and/or Art Unit location of your application in the PTO has changed. To aid in correlating any papers for this application, all further correspondence regarding this application should be directed to Group 180 Art Unit 1807.

Examiner notes that in E.I. du Pont de Nemours & Co. v. Carus Corp., 19 USPQ2d 1174 at 1185 (N.D.Ca. 1991), the court indicated that grant proposals to the NIH and NSF were prior art due to the requirements of the Freedom of Information Act (see 45 C.F.R. §5.1 et seq. and §6.12 et seq.). This may be of some interest to applicants in satisfying 37 C.F.R. 1.56.

Applicants are requested to look over the specification and correct any minor errors.

Restriction to one of the following inventions is required under 35 U.S.C. 121:

I. Claims 1-23, drawn to a method of nucleic acid amplification, classified in Class 435, subclass 6 and 91.

II. Claim 24, drawn to an apparatus and measuring device, classified in Class 435, subclass 291 & 293.

The inventions are distinct, each from the other because of the following reasons: Inventions I and II are related as process and apparatus for its practice. The inventions are distinct if it can be shown that either: (1) the process as claimed can be practiced by another materially different apparatus or by hand, or (2) the apparatus as claimed can be used to practice another and materially different process. (MPEP 806.05(e)). In this case the process as claimed can be practiced by hand as pointed out in the disclosure.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter and different classification, as well as the fact that the search required for Group I is not required for Group II, restriction for examination purposes as indicated is proper.

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During a telephone conversation with Norval B. Galloway on November 4, 1992, a provisional election was made with traverse to prosecute the invention of Group I, claims 1-23. Affirmation of this election must be made by applicant in responding to this Office action. Claim 24 is withdrawn from further consideration by the Examiner, 37 CFR 1.142(b), as being drawn to a non-elected invention. Applicants have indicated that the instant application is a continuation of 07/644,967 which is a continuation of 07/126,920. In both parents, Group I was elected.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a diligently-filed petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(h).

Claims 1-23 are rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 1 and others recite "support capable of specifically associating with the target under binding conditions" which is vague and indefinite functional language describing a chemical moiety by what it does rather than by what it is structurally; therefore it is impossible to know what is and what is not claimed. Claim 6 recites "probe" which is vague and indefinite: do applicants intend a specific nucleic acid sequence which will probe through hybridization or is something else intended? Claim 6 also is phrased in functional language. Claim 10 recites "transcriptase" which is vague and indefinite: was "reverse transcriptase" contemplated? Claim 11 and others recite "non-specific oligonucleotide primer" which is vague and indefinite. Claim 13 and others recite "substantially separating" which is vague and indefinite. Claim 11 recites "capable of binding to a retrievable support" which is vague and indefinite functional language. The claims also recite "retrievable support" but it is not clear what support would not be retrievable thus it is confusing. It also recites "reagents adapted to be applied to said removal product" which is vague and indefinite. Claim 22

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claim 19 since it depends on claim 22) refer to the method of claim 21, but claim 21 is a bit claim corresponding to various compositions of matter. It is not a method claim. This makes claims 22 and 23 confusing. claim 23 recites "capable of interacting with a magnetic field" which is vague and indefinite in light of the known ability of any carbon, nitrogen, or hydrogen containing compound to interact with a magnetic field (e.g. NMR). It is not clear what applicants are describing.

The following is a quotation of 35 U.S.C. 103 which forms the basis for all obviousness rejections set forth in this Office action:

A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Subject matter developed by another person, which qualifies as prior art only under subsection (1) and (2) of section 102 of this title, shall not constitute prior art unless the person whose the subject matter and the claimed invention were, at the time the invention was made, owned or controlled the subject matter, or was subject to an obligation of assignment to the same person.

Claims 1-22 are rejected under 35 U.S.C. 103 as being unpatentable over Mullis when taken with any one of Moss et al., Stabinsky or Engelhardt et al. and taken further in view of Ranki et al. or Josephson or Schroder if necessary.

The primary reference teaches DNA amplification and point out the great value of this method for improved sensitivity as well as improved ability to isolate specific nucleotide sequences. The primary references do not specifically teach nucleic acid affinity chromatography prior to the amplification reaction. The secondary references all teach the well known practice of affinity chromatography, both with nucleic acid attached to a support (surface hybridization) as well as through ligands attached to one strand of the nucleic acid (e.g. biotin-avidin). The secondary references teach the value of affinity chromatography in its ability to isolate specific

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nucleotide sequences and remove unwanted sequences which would interfere with later usefulness of the sequences. The secondary references also teach the greater efficiency of hybridization and improved sensitivity of an already purified sample compared to a non-purified sample (e.g. Moss et al. figure 1) although this fact would be well known to one of ordinary skill in the art. It would be obvious for one of ordinary skill in the art to combine the teachings of the primary references which show improved sensitivity and improved ability to purify a sequence with the secondary references which teach a method providing improved ability to purify a sequence and improved sensitivity since the methods are all directed to the same result and one of ordinary skill would expect an improvement in results.

In regard to claims directed to association with a "probe" it is not clear what applicants mean by this language (see supra); however, it appears to be the well known method of sandwich hybridization (see Rankin et al., this reference has not been provided, it was provided in previous Office Actions in the parent case and it is assumed that applicants are familiar with it) which also claims increased sensitivity and greater ability to isolate specific sequences. In regard to "non-specific oligonucleotide primer" it is not clear what applicants mean by this language (see supra); however, it appears that applicants are simply referring to the well known method of random primer polymerization which is used to label probes. This method is well known not only as an efficient method of making a second copy (into which labeled nucleotides can be added) but is also more efficient than using a single primer. One of ordinary skill in the art would have known this technique and would have been motivated to use it since it makes a second strand thereby doubling the number of copies to be amplified. In regard to the use of a "bead capable of interacting with a magnetic field" it is not clear what applicants mean by this language (see supra); however, it appears to be the well known method of Jahnke and Schneider for magnetic separations. In regard to the kit claims, no prior art references have been provided, they were provided in previous Office Actions on the parent case and it is assumed that applicants are familiar with them. In regard to the kit claims it would have been obvious to one of ordinary skill in the art to package all of the components in a kit for the convenience of practitioners of the method.

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App. filed 12/67

To clarify this rejection, it is examiner's position that applicants simply combined the well known method of nucleic acid amplification with the equally well known method of affinity chromatography to produce a result which could have been expected and with sufficient ingenuity to make the combination. The applicants' invention would have been prima facie obvious at the time of the invention to one of ordinary skill in the art.

Examiner notes that Wood et al., Noyes et al., and Chih et al., which were supplied in previous Office Actions, are merely cumulative to the teachings of Moss et al., Stabinsky and Engelhardt et al., Mullis et al. and Mullis et al. (ref. R) are merely cumulative to Mullis.

No claim is allowed.

This is a continuation of applicant's earlier application SN. 07-04-507. All claims are drawn to the same invention claimed in the earlier application and could have been finally rejected on the grounds or art if rejected in the next Office action if they had been entered in the earlier application. **as a result, THIS ACTION IS MADE FINAL**, even though it is a first action in this case. See MPEP 706.07(b). Applicant is reminded of the provisions of time policy set forth in 37 CFR 1.136(a). The practice of automatically extending the shortened statutory period an additional month upon the filing of a timely first response to a final rejection has been discontinued by the Office. See 1021 TMOG 35.

A SHORTENED STATUTORY PERIOD FOR RESPONSE TO THIS FINAL ACTION IS SET TO EXPIRE THREE MONTHS FROM THE DATE OF THIS ACTION. IN THE EVENT A FIRST RESPONSE IS FILED WITHIN TWO MONTHS OF THE MAILING DATE OF THIS FINAL ACTION AND THE ADVISORY ACTION IS NOT MAILED UNTIL AFTER THE END OF THE THREE-MONTH SHORTENED STATUTORY PERIOD, THEN THE SHORTENED STATUTORY PERIOD WILL EXPIRE ON THE DATE THE ADVISORY ACTION IS MAILED, AND ANY EXTENSION FEE (37 CFR 1.136(a)) WILL BE CALCULATED FROM THE MAILING DATE OF THE ADVISORY ACTION. IN NO EVENT WILL THE STATUTORY

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PERIOD FOR RESPONSE EXPIRE LATER THAN SIX MONTHS FROM THE DATE OF THIS FINAL ACTION.

Examiner has not provided copies of any of the references cited in this Office action because they were provided in earlier Office Actions on the parent case.

An inquiry concerning this communication should be directed to Scott A. Chambers, Ph.D. at telephone number 703-308-3885.

Papers related to this application may be submitted to Group 180 by facsimile transmission. Papers should be faxed to Group 180 via the PTO Fax Center, located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1996 OG 30 (November 15, 1996). The CMI Fax Center number is (703) 306-4227.


Scott A. Chambers
Patent Examiner
Art Unit 1807


MARGARET MOSKOWITZ
SUPERVISORY PATENT EXAMINER
GROUP 180

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FORM PTO-892 (REV. 3-78)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		SERIAL NO. 7/944,505	GROUP/UNIT 1807	ATTACHMENT TO PAPER NUMBER	7
NOTICE OF REFERENCES CITED				APPLICANT(S) Cell			
U.S. PATENT DOCUMENTS							
*	DOCUMENT NO.	DATE	NAME	CLASS	SUB-CLASS	FILING DATE IF APPROPRIATE	
V A	4672040	June 4/87	Josephson	436	526	June 28/85	
V B	4471058	Apr 11/84	Smith et al	436	518		
V C	4486537	Dec 4/84	Rankin et al	436	504		
V D	4687748	Apr 18/87	Schroder	436	526	Mar 25/87	
V E	4751177	June 4/88	Statinaky	435	6	June 3/85	
V F	4683195	Feb 28/87	Mullis et al	435	6	March 28/85	
V G	4683202	Feb 28/87	Mullis	435	91	March 28/85	
H							
I							
J							
K							
FOREIGN PATENT DOCUMENTS							
*	DOCUMENT NO.	DATE	COUNTRY	NAME	CLASS	SUB-CLASS	PERTINENT SHOTS, CWS, ETC.
V L	0097373	04/01/84	EP	Engelhardt	435-6		
M							
N							
O							
P							
Q							
OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, Etc.)							
V R	Mullis et al Cold Spring Harbor Symposium on Quant. Biol. 55 (1980); Cold Spring Harbor Press, CSH N.Y. pp 263-73						
V S	Moss et al Journal of Biol. Chem 256:12655-58(1981)						
V S	Shih et al Biochemistry 13(16):3411-18(1974)						
V T	Foyes et al., Cell 5 301-11 (1975)						
V U	Wood et al., Journal of Biol Chem 252:457-63 1977						
EXAMINER		DATE					
J. H. Schumacher		11/4/92					

* A copy of this reference is not being furnished with this office action.
(See Manual of Patent Examining Procedure, section 707.05 (a).)

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GROUP

APPLICATION NUMBER

9441505

NOTICE OF DRAFTSMAN'S PATENT DRAWING REVIEW

The PTO Draftsmen review all originally filed drawings regardless of whether they were designated as informal or formal.

The drawings that:

A ☐ are approved.B ☒ are rejected or corrected. The drawings checked below. This examination requires submission of new corrected drawings at the appropriate time. Corrected drawings must be submitted according to the instructions listed on the back of this Notice.

1. Paper and Ink. 37 CFR 1.84(a)

☐ Poor Quality Paper Must Be White.☐ Transparent Paper Not Allowed.

Fig(s):

2. Size of Sheet and Margins. 37 CFR 1.84(b)

Acceptable Paper Sizes and Margins

Paper Size

Margin	14 inches	12 inches	11 inches by 21 by 29.7 cm.
Top	2 inches	1 inch	2.5 cm.
Left	1/4 inch	1/4 inch	2.5 cm.
Right	1/4 inch	1/4 inch	1.5 cm.
Bottom	1/4 inch	1/4 inch	1.0 cm.

☐ Proper Size Paper Included in Sheets Must Be Same Size.

Fig(s):

☒ Proper Margins Required

Sheet(s) 6-8

☒ Top ☐ Right
☐ Left ☐ Bottom

3. Character of Lines. 37 CFR 1.84(c)

☒ Lines Pale, Rough and Blurred, or Jagged. Fig(s) 4-6☐ Solid Black Shading Not Allowed. Fig(s):4. ☐ Photographs Not Approved.☐ Comments:

5. Hatching and Shading. 37 CFR 1.84(j)

☐ Cross Hatching Required.☐ Hatching Lines Must Be Parallel.

Fig(s):

☐ Double Line Hatching Not Allowed.

Fig(s):

☐ Parts in Section Must Be Hatched Properly. Fig(s):

6. Reference Characters. 37 CFR 1.84(f)

☒ Reference Characters Poor or Rough and Blurred. Fig(s) 4-6

The number 1/8 inch (3.2 mm.) in height

☒ Reference Characters Poor or Rough and Blurred. Fig(s) 4-7

7. Views. 37 CFR 1.84(g) & (h)

☐ Figures Must Be Numbered Separately.☐ Figures Must Not be Connected. Fig(s):

8. Identification of Drawings. 37 CFR 1.84(i)

☐ Extraneous Matter or Copy Machine Marks Not Allowed. Fig(s):9. ☐ Changes Not Completed from Prior PTO-948 dated:

Telephone inquiries concerning this review should be directed to the Chief Draftsman at telephone number (703) 557-6404.

Reviewing Draftsman

Date